

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

PATENT SPECIFICATION (11)

1 571 832

1 571 832

- (21) Application No. 253/77 (22) Filed 5 Jan. 1977 (19)
 (31) Convention Application No. 661612 (32) Filed 26 Feb. 1976 in
 (33) United States of America (US)
 (44) Complete Specification Published 23 Jul. 1980
 (51) INT. CL.³ A61K 9/06
 (52) Index at Acceptance
 A5B 826 L



(54) MEDICAL DRESSINGS CONTAINING PHARMACEUTICALLY ACTIVE MATERIAL

(71) We, FLOW PHARMACEUTICALS, INC., a corporation of the State of Nevada, United States of America, of 3780 Fabian Way, Palo Alto, California, United States of America, do hereby declare the invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:-

This invention relates to a new medical dressing carrying a pharmaceutically active material, that is a drug or medicament, and more particularly to a medical dressing formed *in situ* on a mucous membrane from a pharmaceutical composition which is liquid at ambient temperatures and which gels on the mucous membrane owing to the elevated temperature possessed thereby. The pharmaceutically active material is supplied directly to the site to be treated.

There are many drugs known to be useful for treatment of afflictions or protection of mucous membranes, e.g., for ocular diseases. A practical problem in connection with therapeutic or protective application of pharmaceutically active chemicals to afflicted mucous membranes resides in the delivery of the chemical or drug to the affected area in need of treatment. Various formulations and techniques have been attempted to deliver medicaments to mucous membranes, but there is a need for improved pharmaceutical vehicles for delivery of drugs, and it is this need to which the present invention is addressed.

For example, drugs have been formulated into aqueous solutions. However, the fleeting presence and poor contact of aqueous solutions applied to mucous membranes has been a disadvantage. The only adequate application of medication in solution form to mucous membranes is usually accomplished by employing continuous lavage or interrupted irrigation. This approach is often wasteful of expensive drugs and poses a major problem of inconvenience. Thus, treatment of severe keratoconjunctivitis sicca with isotonic salt solutions requires ocular instillation every 15-30 minutes.

The use of viscous aqueous solutions is usually more convenient. For example, the aforementioned isotonic salt solutions can often be applied every 1-2 hours and accomplish the same therapeutic objective if the solution is made viscous. Drugs have been formulated into aqueous suspensions made viscous by the addition of gums or cellulose-modified synthetic derivatives or incorporated into oleaginous vehicles or bases consisting of natural plant or animal fats or modifications thereof or petroleum-derived hydrocarbons.

Indeed, aqueous vehicles which are thickened by the addition of selected gums or cellulose-derived viscosity building agents are perhaps the most commonly used media for delivery of drugs or medicaments to mucous membranes. Generally, the viscosity of such preparation ranges from about 25 cps to indeterminate values in stiff gels. Nearly uniform drug delivery is possible with such vehicles, and they frequently provide desirable protection to the mucous membranes.

In contrast, non-viscous aqueous suspensions have many disadvantages and are not typically used. A major problem is rapid settling of the suspended drug. This gives rise to undesirable need for continuous stirring during administration in order to deliver a uniform dose.

While thick gels would seem to offer the best potential in terms of protection as well as holding and delivering medication, they in fact have some disadvantages. In some instances, they are difficult to apply from their respective commercial containers. Moreover, thick gels do not spread readily over the area being treated, and possibly painful spreading and rubbing may be necessary. Also, on evaporation of the water from the vehicle, a cosmetically

2

1,571,832

2

unappealing hard granular or flaky residue often results at the site of the application.

Attempts to use oily vehicles to increase drug delivery and prolong ensuing pharmacologic action have not met with uniform success. The use of oleaginous vehicles, whether anhydrous or in emulsion form (oil-in-water or water-in-oil), may have advantages for certain therapeutic indications, if the vehicle will adhere. However, since normal mucous membranes are always moist with aqueous tissue fluids, and water does not mix readily with oil bases, application, uniform spreading, and retention all become difficult. Perhaps the only time oily or emulsion vehicles are used successfully is when the mucous tissue is abnormally dry because of disease.

Another approach to the delivery of drugs or medicaments to mucous membranes is the recent development of silicone plastic devices which deliver drugs at predetermined, nearly uniform, zero order rates extending from a few days to several years. However, the usefulness of such devices depends upon a constant supply of tissue fluid or glandular secretion; in the absence of fluid, plastic devices are not operative. Such devices are not designed to offer any protection to an inflamed mucous membrane. Discomfort often associated with the devices, and inadvertent loss of the devices, are additional problems.

The existence of all these disparate approaches to drug delivery to mucous membrane evidences the need for the new pharmaceutical vehicles. Against the background of this array of formulations and devices with all their attendant problems, the present invention fills that need.

According to the present invention, there is provided a medical dressing applied to mucous membrane, which dressing comprises a gelled pharmaceutical composition conforming at least in part to the configuration of the mucous membrane, which pharmaceutical composition has been applied to the mucous membrane at a temperature below human body temperature and at which it is flowable, which pharmaceutical composition consists of a clear water-miscible physiologically acceptable liquid vehicle which is flowable at ambient temperatures and which gels to a thickened non-flowing and adhering consistency at human body temperature, the liquid vehicle comprising an aqueous solution of a polyoxyethylene polyoxypropylene block copolymer and having a gel transition temperature in the range of from 25° to 40°C, which aqueous solution contains therein a pharmacologically effective amount of a pharmacologically active chemical material.

The pharmaceutical composition from which the dressing is produced contains a pharmaceutical vehicles which is liquid at ambient temperatures below 30°C, preferably below 25°C. It has a sol-gel transition temperature usually in the range of from 25°C to 40°C, preferably from 25°C to 35°C and most preferably from 29°C to 31°C.

More particularly, it has been discovered that aqueous solutions of certain polyoxyethylene-polyoxypropylene block copolymers are useful pharmaceutical vehicles having the properties set forth above. Preferred polyoxyethylene-polyoxypropylene block copolymers for use in compositions of the present invention are polyoxyethylene-polyoxypropylene block copolymers in which the number of polyoxyethylene units is at least 50%, preferably approximately 70% of the total number of monomeric units in the total molecule, the block copolymers having an average molecular weight of from 7500 to 15,500, preferably approximately 11,500.

The block copolymers preferably constitute:- from 10% to 26% more preferably from 12% to 26%, by weight of the pharmaceutical composition, the composition containing preferably from 74% to 90% more preferably from 74% to 83% by weight water. The block copolymers preferably possess a room temperature solubility in water of greater than 10 grams per 100 ml. of water, and a cloud point in 1% aqueous solution of at least 100°C.

The pharmaceutical composition may also include various additives, such as auxiliary non-ionic surfactants, salts to adjust osmotic pressure, buffer systems to control pH, and preservatives. Preferably, the composition contains at least one water-soluble compatible salt for adjustment of osmotic pressure in sufficient amount to provide a solution salt content equivalent to up to 10%, usually from 0.1% to 10.0%, and preferably from 0.5% to 6.0% sodium chloride. It is also preferred that the composition contain a compatible preservative or germicide in an amount effective to afford protection to the vehicle against bacterial contamination.

In accordance with this invention, the pharmaceutical composition preferably has a pH in the range of 3.5 to 9.5. Particularly preferred is a pH in the range of from 6.0 to 8.5, and especially from 6.2 to 7.8.

The concept of this invention is not dependent on the nature of the pharmaceutically active material or drug and any compatible pharmaceutically active material or drug may be used. Preferably, the drug is water-soluble. However, drugs which are not ordinarily soluble in water may also be employed, and where needed, auxiliary nonionic surfactants, which are typically well tolerated by mucous membranes, can be added to increase the solvent action, while maintaining the vehicle gel transition temperature within the required range.

3

1,571,832

3

It has been discovered that a wide variety of useful pharmaceuticals which are not ordinarily soluble in water and are presently marketed only in suspension form can in fact be dissolved in the polyoxyethylene-polyoxypropylene vehicles of the present invention. In some instances, the addition of auxiliary nonionic surfactants was found necessary. However, the critical gel transition temperature is maintained.

An important aspect of this invention is that the pharmaceutical compositions are liquid at ambient room temperatures and can be applied to the affected mucous membrane area by conventional liquid depositing means, including dispensation to the area of treatment from standard plastic squeeze bottles or in drop form. At body temperatures above 30°C the vehicle or base passes through the sol-gel transition temperature and gels to a thickened, non-flowing and adhering consistency, thereby to form a medical dressing on the mucous membrane holding and delivering the medication as required and for prolonged periods of time.

As previously indicated herein the pharmaceutical vehicle of a pharmaceutical composition used to form a medical dressing of this invention consists of a clear, water-miscible, physiologically-acceptable medium which is liquid at ambient temperature below about 30°C. and thickens to a gel at body temperatures above about 30°C. In practice, it has been found that a vehicle having a sol-gel transition temperature in the range of from 25°C to 40°C. satisfies this requirement and is useful in the practice of the present invention. Preferably, the sol-gel transition temperature will be in a range of from 25°C to 35°C, and excellent results have been obtained using vehicles having a sol-gel transition temperature in the range of from 29 to 31°C.

The capacity of the liquid pharmaceutical vehicle to gel at human body temperatures is the critical feature of the invention for it is in this property that many of the disadvantages of the previous approaches are overcome. This, the dissipative quality of aqueous solutions is avoided since the vehicles herein gel at the site of treatment. Moreover, the problems of formulation, handling and application of viscous aqueous vehicles or gels are overcome since at the time of application the present pharmaceutical vehicle and composition are free-flowing liquids.

The pharmaceutical vehicle employed in compositions used to form medical dressings of this invention is clear and water-miscible. These are especially important requirements for usefulness in therapeutic and protective ocular applications. Water-miscibility of the vehicle overcomes major problems faced in attempts to use oily vehicles.

The vehicle of must be physiologically acceptable so that no adverse reaction occurs when the pharmaceutical composition comes in contact with human tissue or fluids. Thus, the vehicles must be inert when tested for ocular tolerance in human and rabbit eyes;

As also previously indicated herein, a suitable pharmaceutical vehicle comprises an aqueous solution of a selected polyoxyethylene-polyoxypropylene block copolymer. It has been found that polyoxyethylene-polyoxypropylene block copolymers in which the number of polyoxyethylene units is at least 50% of the number of units in the total molecule, the block copolymer having an average molecular weight of from 7500 to 15,500 a room temperature solubility in water, preferably greater than 10 grams per 100 ml. of water, and a cloud point in 1% aqueous solution preferably of at least about 100°C., can be used to form composition having a sol-gel transition temperature in the range of from 25°C to 40°C.

Such block copolymers are included in a series of nonionic surface-active agents sold under the trademark "Pluronic" by Wyandotte Chemical Corp. The "Pluronics" are closely related block copolymers that may be generically classified as polyoxypropylene-polyoxyethylene condensates terminating in primary hydroxyl groups. They are formed by the condensation of propylene oxide into a propylene glycol nucleus followed by the condensation of ethylene oxide onto both ends of the polyoxypropylene base. The polyoxyethylene hydrophilic groups on the ends of the molecule are controlled in length to constitute anywhere from 10% to 80% by weight of the final molecule.

The "Pluronic" series of products may be represented empirically by the formula:



where a and c are statistically equal. They have been available in average molecular weights of from about 1100 to 15,500.

A preferred polyoxyethylene-polyoxypropylene block copolymer for use in the pharmaceutical composition is one in which the number of polyoxyethylene units is approximately 70% of the total number of monomeric units in the molecule and where the copolymer has an average molecular weight approximately 11,500. "Pluronic F-127" is such a material, and it has a solubility greater than 10 gms/100 ml. water as well as a cloud point in 1% aqueous solution higher than 100°C (Pluronic is a Registered Trade Mark).

1.571,832

4

The concentration of the polyoxyethylene-polyoxypropylene condensate is an important parameter. Significantly, by ready adjustment of the concentration of the copolymer to accommodate other solutes present in the vehicle, any desired gel transition temperature in the critical range of above ambient temperature and below body temperature can be achieved. Thus, the principal consideration is the selection of a concentration which, in conjunction with all of the constituents of the vehicle composition, will provide a sol-gel transition temperature in the required range.

5

It has been found that a useful block copolymer concentration is from 10% to 26% by weight, particularly from 17% to 26%. Excellent results have been obtained using aqueous solutions of from 17% to 26% by weight of "Pluronic F-127". The water content is generally from 74% to 90% by weight of the vehicle composition, and is typically from 74 to 85% by weight. The water used in forming the aqueous solution is preferably purified, as by distillation, filtration, ion-exchange or the like.

10

The polyoxyethylene-polyoxypropylene pharmaceutical vehicles used in compositions of this invention have been unexpectedly found to increase drug absorption by the mucous membrane. Moreover, it has also been found that the pharmacologic response is unexpectedly prolonged. Drug action is typically both increased and prolonged by a factor of 2 or more. At the same time, protection is afforded to the involved tissues.

15

Another advantage is that they are compatible with the therapeutic bandage semi-hard (silicone) and soft or flexible contact lenses. In contrast to drug suspensions in which suspended particles could be lodged in the surfaces of the lenses and cause focal points of irritation or blurred vision, and in contrast to oily vehicles or bases which could adversely affect lens clarity, degree of hydration, and the physical parameters of therapeutic lenses, the present vehicles, when used in conjunction with therapeutic contact lenses, markedly increased wearing comfort, provided cleaner lenses, and gave more rapid healing responses than without instillation of the vehicle.

20

The liquid pharmaceutical compositions from which the dressings of this invention are produced preferably include at least one water-soluble compatible salt to adjust osmotic pressure. Frequently, the composition will be formulated to be isotonic with human serum and tear fluid, the normal tonicity of which is 0.9% (9.0 grams of sodium chloride) per litre of the composition. Isotonic solutions contain about 0.9% sodium chloride, or other salt or mixture of salts having a salt content equivalent to about 0.9% sodium chloride in their osmotic effect.

25

In general, the compositions may contain a sufficient amount of at least one salt to provide up to about 10%, especially from 0.5% to 6.0%, sodium chloride equivalent salt content. Polyoxyethylene-polyoxypropylene vehicles with as high as 10% sodium chloride equivalent salt content can be made having the requisite gel transition temperature. Such compositions are markedly hypertonic, and can be advantageously used where commercially available hypertonic solutions are presently employed.

30

Generally, it was found that each additional increment of salt proportionately lowered the gel transition temperature.

35

Any soluble salt or mixture of salts compatible with mucous membrane tissue can be used to provide the desired tonicity. Sodium chloride, potassium chloride, or mixtures thereof, are presently preferred. However, one or more essentially neutral, water soluble alkali metal salts can be substituted in whole or in part for the sodium or potassium chloride in the vehicles of this invention. Thus, other alkali metal halides, such as sodium bromide, potassium fluoride or potassium bromide can be used. Other salts, such as sodium sulfate, potassium sulfate, sodium nitrate, sodium phosphate, potassium nitrate or potassium phosphate can also be used.

40

Preferably, the pharmaceutical composition contains a compatible preservative or germicide in an amount effective to afford protection to the vehicle against bacterial contamination. Any conventional preservative system may be used.

45

Quaternary germicides, particularly benzalkonium chloride, are presently preferred. Benzalkonium chloride is an alkyl substituted dimethylbenzylammonium chloride in which the alkyl substituents comprise a mixture of C₈ to C₁₂ alkyl radicals. Exemplary of other preservatives which can be desirably used are salts of ethylenediaminetetracetic acid, known as edetates, such as disodium edetate and trisodium edetate, sorbic acid, salts of sorbic acid, boric acid, and salts of boric acid, such as sodium borate. Still other useful preservatives or germicides are thimerosal sodium, phenylmercuric acetate, methyl, ethyl and propyl para-aminobenzoic acid esters.

50

The preservatives can be used individually or in combination. They are used in effective amount to afford protection against contamination. For example, amounts of from 0.001% to 0.03% by weight of a quaternary or organic mercurial germicide are known to be effective and can be used in the present invention. Sorbic acid NF XIII is known to be useful in amounts of from 0.01% to 0.5% by weight and may be so used in the pharmaceutical

55

60

65

5

1,571,832

5

compositions.

The pH of the pharmaceutical compositions may be adjusted as desired. In general, the pH can range from 3.5 to 9.5. Preferably, the pH is from 6.0 to 8.5, and especially from 6.2 to 7.8, the range of the human tear. In some instances, the stability of certain preservatives is maximized by pH adjustment. For example, acid to neutral pH is optimal for the alkyl para-amino-benzoic acid esters.

Compatible, conventional buffers, i.e., weak acids, weak bases, and their corresponding salts, may be used to adjust pH as desired. A sodium biphosphate, disodium phosphate system is exemplary of useful buffering systems. An effective amount of buffer is used to achieve the desired pH. For example, a combination of from 0.2% to 0.6% sodium biphosphate and from 0.2% to 0.7% disodium phosphate may be used to adjust to a pH in the 6.2 to 7.2 range. Certain preservatives also affect pH, such as trisodium edetate. By selection and simple correlation of the desired additives, one having ordinary skill in the art can readily adjust the pH as desired, while retaining the gel transition temperature in the required range.

Compatible and physiologically-acceptable auxiliary nonionic surfactants may optionally be used to improve solvation of the drug or medicament. Exemplary of conventional surfactants which may be used are Polysorbate 80 and polyoxyl 40-stearate employed in conventional amounts.

Any pharmaceutically active material may be admixed in a pharmacologically effective amount with the pharmaceutical vehicle to form the pharmaceutical compositions of this invention. Preferably, the drug is water-soluble. However, drugs which are not ordinarily soluble in water may also be employed, and it has been found that a wide variety of useful drugs which are currently marketed in suspension form can be dissolved in the polyoxyethylene-polyoxypropylene vehicles of the present invention. Where necessary or desirable, auxiliary nonionic surfactants may be included in the pharmaceutical composition.

The drug or medicament is selected on the basis of the treatment indicated for the patient. Exemplary of drugs which have been used in connection with the pharmaceutical vehicles herein are pilocarpine HCl for glaucoma, phylephrine for red eyes and Dexamethasone U.S.P., for inflammatory ocular conditions. Various anti-microbial pharmaceuticals for treatment of fungal and viral diseases of mucous membranes may be used, such as Clofazimine, pimaricin, amphotericin, neomycin sulfate, choramphenical, bacitracin, sulfacetamide, gentamycin and polymix in B sulfate.

The pharmaceutical used to produce the medical dressings compositions of this invention can be readily prepared. Essentially, any solution forming technique may be used. The vehicle may be prepared separately and the pharmaceutical added thereto, or preferably, the pharmaceutical composition is formulated without separate preparation of the vehicle. For example, in the use of the polyoxyethylene-polyoxypropylene block copolymer vehicles, the pharmaceutical composition is desirably prepared by fusing the block copolymer, adding the pharmaceutically active material to the fused copolymer, and dissolving the pharmaceutical by simple stirring. A water solution of the remaining ingredients is prepared, and the solution of pharmaceutical in the block copolymer is mixed with the aqueous solution to form a solution of all components. The pH may then be adjusted as desired, e.g., by addition of a basic or acidic solution as desired. It is generally preferred to add copolymer or a solution of a pharmaceutically active material in the copolymer to the water or aqueous solution rather than adding the water or aqueous solution to the copolymer or copolymerpharmaceutical mixture.

The pharmaceutical composition is a liquid at ambient temperatures and therefore may be employed in any manner conventionally used to apply free-flowing liquid pharmaceuticals to mucous membranes. Preferably, application is in drop form in the manner typically used, for example, to apply eye drops. Thus, the normal squeeze-type liquid drop application devices are perfectly suitable for use in applying the pharmaceutical compositions of this invention to the site intended for treatment. The amount of pharmaceutical composition should be sufficient to deliver a pharmacologically effective amount of the active pharmaceutical to the mucous membrane treatment area.

In addition to overcoming major disadvantages of previous techniques for delivering drugs and medicaments to mucous membranes, since a medical dressing is formed *in situ*, the present invention has been found to increase drug absorption by the affected tissue and prolong pharmacologic response. Many other advantages will be apparent to those skilled in the art.

The following examples illustrate the compositions of the present invention, and their preparation and utility, but are not limitative of the invention. All percentages are standard weight in volume (W/V) % expressions. In each instance, the formulations were made sterile by using standard heat and pressure techniques, as well as aseptic techniques.

Example I

A pharmaceutical vehicle for use in forming a medical dressing according to this invention

6

1,571.832

6

is exemplified by the composition:

	Pluronic F-127	18%
	sodium chloride	0.75%
5	potassium chloride	0.25%
	disodium edetate	0.025%
	benzalkonium chloride	0.004%
	purified water, enough to make	100%
10	(adjust pH to 7.4 with dilute sodium hydroxide solution)	

5

10

It is most easily prepared by mixing all the ingredients in 95% of the required water and allowing the polyoxyethylene-polyoxypropylene copolymer (Pluronic F-127) to hydrate and completely dissolve overnight with gentle stirring at temperatures below 20°C. Once a clear solution is obtained, the pH is adjusted to pH 7.4; and the balance of the water is added. The sol-gel transition temperature was found to be at 29-30°C.

15

15

This vehicle formulation was evaluated on rabbit eyes according to the Draize scoring technique. On a scale having a maximum of 110 possible units of irritation or ocular trauma, experimental values were consistently at or near zero indicating that it produces no adverse ocular effects.

20

20

The vehicle itself has pharmacologic utility. It was tested for use in alleviating ocular symptoms of Sjogren's syndrome. Two ophthalmologists treated 11 patients and reported that almost instant relief was obtained.

Example II

25

25

A variation of the Example I composition was prepared for testing in bullous keratopathy as follows:

	pluronic F-127	17	%
30	benzalkonium chloride	0.0075%	
	disodium edetate	0.0125%	
	trisodium edetate	0.025	%
	sodium chloride	3.75	%
35	potassium chloride	1.0	%
	purified water, enough to make	100%	

30

35

The desired pH is achieved and maintained by the use of the acidic and basic salts of ethylenediaminetetraacetic acid. This formulation contains approximately five times the amount of salt present in isotonic sodium chloride solution. The solgel transition temperature was about 30°C.

40

40

When the formulation was tried in 6 patients, with and without soft contact lenses, and all with corneal edema associated with bullous keratopathy, very significant improvement was noted. In several other instances where corneal swelling was associated with prolonged hard contact lens wear, as well as other edematous conditions of unrelated problems the product was found efficacious. When compared to Adsorbonac S (Burton Parsons, Inc, Washington, D.C.), a commercially available hypertonic solution, the new invention was preferred in all instances. The test preparation caused significantly less discomfort upon eye instillation.

45

45

Example III

50

50

The following vehicle was prepared:

	Pluronic F-127	16	%
	sorbic acid	0.1	%
	disodium edetate	0.1	%
55	sodium borate	0.23%	
	sodium chloride	0.5	%
	potassium chloride	0.2	%
60	purified water, enough to make	100%	

55

60

The pH was 7.5, and the sol-gel transition temperature was 34°C. With the advent of continuous wear therapeutic soft contact lenses, and more recently continuous wear cosmetic lenses, there is a frequent need for innocuous eye drops that loosen the accumulated mucoid deposits on the lenses, reequilibrate the lenses and add to the overall comfort of wear.

65

65

7

1,571,832

7

The composition of this Example was evaluated by an ophthalmologist for this purpose by eye drop application on 14 patients. Without exception, all of the patients found that this eye drop was the best product which they had used. The ophthalmologist was also impressed with the clinical response.

Examples I, II and III illustrate compositions which may be used as pharmaceutical vehicles when producing medical dressings in accordance with the invention, and significantly, have protective and therapeutic usefulness in themselves without further addition of drugs or medicaments. The following Examples demonstrate preparation and utility of pharmaceutical compositions in accordance with the invention.

Example IV

The following pharmaceutical composition containing the drug Dexamethasone as the added active pharmaceutical material was prepared to test corticosteroid anti-inflammatory solubilization and stability:

Dexamethasone U.S.P.	0.05 %
Pluronic F-127	19 %
thimerosal sodium U.S.P.	0.005 %
disodium edetate U.S.P.	0.1 %
sodium chloride U.S.P.	0.9 %
purified water U.S.P., enough to make 100%	

To prepare this formulation, the Pluronic F-127 was first fused at about 50-60°C., at which point the Dexamethasone was added and dissolved by simple stirring. The remainder of the ingredients were dissolved in water and added. Then the pH was adjusted to 7.0 with a dilute solution of sodium hydroxide. The solution, observed over a period of 5 months, remained crystal clear. The sol-gel transition temperature was about 26°C. On warm days, refrigeration was required to maintain the product in the liquid state. However, this turned out to be advantageous for when the cooled product was tried on 2 patients with a severe inflammatory ocular condition resulting from chemical burns, the cooling sensation upon instillation provided added relief. Other clinical tests of this formulation verified its utility and demonstrated that the concentration herein employed, which is half of the amount normally used in the commercially available product Decadron (Registered Trade Mark of Merck, Sharpe & Dohme), was at least as effective. Using half as much of an expensive raw material could mean a significant saving to the patient.

Example V

The following pharmaceutical composition containing pilocarpine HCl as the added active pharmaceutical was prepared for treatment of glaucoma:

Pluronic F-127	18 %
sodium chloride	0.3 %
potassium chloride	0.1 %
disodium phosphate	0.5 %
sodium biphosphate	0.08 %
benzalkonium chloride	0.01 %
purified water, enough to make 100%	
pH - 6.8	

This formulation was compared to 2% pilocarpine (commercially available in an aqueous solution) normally prescribed for glaucoma in 4 patients. In all instances, the reduction in intra-ocular pressure by treatment with the present formulation was found to be as good as or better than the product having four times the concentration.

Example VI

The following composition containing phenylephrine HCl as the added pharmaceutically active material was prepared:

phenylephrine HCl	0.1 %
Pluronic F-127	18 %
sodium chloride	0.9 %
benzalkonium chlorid	0.008 %
purified water, enough to make 100%	

8

1,571,832

8

This formulation was compared to an aqueous 0.5% solution in a small series of patients with red eyes. The rate of vasoconstriction (scleral blanching) in both instances was about the same. Two of the 3 volunteer patients reported better comfort in the eye treated with the present formulation. In all 3 patients, the paired eyes treated with this product looked much better than the 0.5% phenylephrine solution when examined with a slit lamp 20 minutes after treatment. Residual amounts of the pharmaceutical vehicle were still apparent in the treated eyes, whereas all of the more concentrated 0.5% aqueous solution had dissipated in the opposite eyes. This observation demonstrates the added ocular protection and duration of the new drug form of this invention.

Example VII

The following pharmaceutical solution containing the antimicrobial agent Clofazimine was prepared:

Clofazimine	0.1%
Pluronic F-127	12 %
Polysorbate 80	20 %
sodium chloride	0.6%
benzalkonium chloride	0.1%
purified water, enough to make 100%	
pH - 6.8, sol-gel transition	
at 35°C.	

This pharmaceutical composition was tested in vitro and found to exhibit good activity.

Example VIII

The following pharmaceutical solution containing the antimicrobial agent pimaricin was prepared:

pimaricin	0.3%
Pluronic F-125	10 %
(average molecular weight of about 8000, polyoxyethylene units about 50% of total units in molecule)	
polyoxyl 40-stearate	20 %
sodium chloride	0.6%
benzalkonium chloride	0.1%
purified water enough to make 100%	
pH - 6.5, sol-gel transition	
at 31°C.	

This formulation was also tested in vitro and was likewise found to exhibit good activity.

Pharmaceutical compositions containing antimicrobial agents other than those of Examples VII and VIII have similarly been prepared and tested with success. Suitable vehicles for antimicrobial agents have been a recognized problem, and the usefulness of the vehicles of this invention in connection with antimicrobial agents represents a particularly significant and advantageous aspect of this invention.

WHAT WE CLAIM IS:-

1. A medical dressing applied to mucous membrane, which dressing comprises a gelled pharmaceutical composition conforming at least in part to the configuration of the mucous membrane, which pharmaceutical composition has been applied to the mucous membrane at a temperature below human body temperature and at which it is flowable, which pharmaceutical composition consists of a clear water-miscible physiologically acceptable liquid vehicle which is flowable at ambient temperatures and which gels to a thickened non-flowing and adhering consistency at human body temperature, the liquid vehicle comprising an aqueous solution of a polyoxyethylene - polyoxypropylene block copolymer and having a gel transition temperature in the range of from 25° to 40°C, which aqueous solution contains therein a pharmacologically effective amount of a pharmacologically active chemical material.

2. A dressing according to claim 1, in which said liquid vehicle is an aqueous solution containing from 10% to 26% by weight of a polyoxyethylene-polyoxypropylene block copolymer in which the number of polyoxyethylene units is at least 50% of the total number of units in the copolymer, the copolymer having an average molecular weight of from 7500 to 15,500.

3. A dressing according to claim 2, in which the copolymer constitutes from 12% to 26% by weight thereof, has an average molecular weight of approximately 11,500, and in which

the number of polyoxyethylene units is approximately 70% of the total molecule, the dressing having been formed by gelling of the liquid at a temperature from 25 to 35°C.

4. A dressing according to any one of the claims 1 to 3, which contains a sufficient amount of at least one water-soluble compatible salt to provide a solution salt content equivalent to up to 10% sodium chloride.

5. A dressing according to claim 4, wherein the salt is selected from sodium halide, sodium sulfate, sodium nitrate, sodium phosphate, potassium halide, potassium sulfate, potassium nitrate, potassium phosphate and mixtures thereof.

6. A dressing according to claim 4 or 5, in which the solution salt content is equivalent to from 0.5% to 6.0% sodium chloride.

7. A dressing according to any one of the claims 1 to 6, which contains a compatible preservative in an amount effective to afford protection to the dressing against bacterial contamination.

8. A dressing according to claim 7, wherein the preservative is selected from benzalkonium chlorides, a sodium salt of ethylenediaminetetraacetic acid, sorbic acid, boric acid an alkali metal salt of boric acid, thimerosal sodium, phenylmercuric acetate, an alkyl ester of para-aminobenzoic acid, and mixtures thereof.

9. A dressing according to any one of the claims 1 to 8, in which the pharmaceutically effective chemical material is selected from dexamethasone, pilocarpine HCl, phenylephrine HCl and an antimicrobial agent.

10. A dressing according to any one of claims 1 to 9, which has a pH of from 6.0 to 8.5.

11. A dressing as claimed in any one of the preceding claims, which is applied to ocular mucous membrane.

12. A medical dressing as claimed in claim 14, substantially as described in any one of the foregoing examples IV to VIII.

HASELTINE, LAKE & CO.

Chartered Patent Agents.

Hazlitt House,

28, Southampton Buildings,

Chancery Lane, London WC2A 1AT

-also-

Temple Gate House, Temple Gate, Bristol BS1 6PT

-and-

9, Park Square, Leeds, LS1 2LH, Yorks